

SEQUENCE LISTING

**GENERAL INFORMATION:** 

(i)

APPLICANT: PEREGRINO FERREIRA, Paulo;

5 GESSIEN KROON, Erna;

PIMENTA DOS REIS, Karlisson Jennner;

BIAS FORTES FERRAZ, Isabella;

CERQUEIRA LEITE, Romulo.

(ii)

TITLE OF INVENTION: Method and composition for the diagnosis of equine infectious anemia virus disease by using the recombinant capsid protein virus (p26)

(iii)

**NUMBER OF SEQUENCES: 1** 

15 (iv)

**CORRESPONDENCE ADDRESS:** 

(A)

ADDRESSEE: Universidade Federal de Minas Gerais - CTIT

(B)

20 STREET: Avenida Antônio Carlos, 6627 Bairro São Francisco

(C)

CITY: Belo Horizonte

(D)

STATE: Minas Gerais

25 **(E)** 

**COUNTRY: BRAZIL** 

(F)

ZIP: 31270-901

(v)

30 COMPUTER READABLE FORM:

(A)

MEDIUM TYPE: diskette - 3.50 inch, 1.44 Mb storage

(B)

COMPUTER: IBM compatible

(C)

5 OPERATING SYSTEM: Windows 98

(D)

SOFTWARE: Office premium

(vi)

**CURRENT APPLICATION DATA:** 

10 **(A)** 

APPLICATION NUMBER: U.S. 09/331.262

(B)

FILING DATE:

(C)

15 CLASSIFICATION: C12Q1/70

(vii)

PRIOR APPLICATION DATA

(A)

APPLICATION NUMBER: PI 9606273-8

20 (B

FILING DATE: 18-DEC-1996

(2)

**INFORMATION FOR SEQ ID N0:1:** 

(i)

25 SEQUENCE CHARACTERISTICS:

(A)

LENGHT: 252 amino acids

(B)

TYPE: amino acid

30 **(D)** 

TOPOLOGY: linear

```
(ii)
    MOLECULE TYPE : protein
    (vi)
    ORIGINAL SOURCE
    (A)
    ORGANISM: equine infectious anemia virus
    (ix)
    FEATURE:
    (A)
10
    NAME: p26
    (x)
    PUBLICATION INFORMATION
    (A)
    AUTHORS:
15
    (B)
    TITLE: (
    C)
    JOURNAL:
    (D)
20
    VOLUME:
    (F)
    PAGES:
    (G)
    DATE:
25
    (xi)
    SEQUENCE DESCRIPTION: SEQ ID NO:1
    His His His His His Gly Ser Pro Gly Asn Pro Leu Thr Trp
                  5
                                   10
                                                     15
30
```

	Ser Lys Ala Leu Lys Lys Leu Glu Lys Val Thr Val Gln Gly Ser				
	20	25	30		
	Gln Lys Leu Thr Thr Gly Asn Cys Na Trp Ala Leu Ser Leu Val				
	35	40	45		
5	Asp Leu Phe His Asp Thr	Asn Phe Val Lys Glu Lys	Asp Trp Gln		
	50	55	60		
	Leu Arg Asp Val Ile Pro Leu Leu Glu Asp Val Thr Gln Thr Val				
	65	70	75		
	Ser Gly Gln Glu Arg Glu A	la Phe Glu Arg Thr Trp T	rp Ala IIe		
10	80	85	90		
	Ser Ala Val Lys Met Gly Leu Gln lle Asn AsnVal Val Asp Gly				
	95	100	105		
	Lys Ala Ser Phe Gln Leu L	∟eu Arg Ala Lys Tyr Glu L	ys Lys Thr		
	110	115	120		
15	Ala Asn Lys Lys Gln Ser G	Blu Pro Ser Glu Glu Tyr P	ro Ile Met		
	125	130	135		
	lle Asp Gly Ala Gly Asn Ar	rg Asn Phe Arg Pro Leu T	hr Pro Arg		
	140	145	150		
	Gly Tyr Thr Thr Trp Val As	snThr lle Gln Thr Asn Gly			
20	155	160	165		
	Asn Glu Ala Ser Gln Asn I	_eu Phe Gly Ile Leu Ser V			
	170	175	180		
	Thr Ser Glu Glu Met Asn A	·	-		
	185	190	195		
25	Ala Gly Gln Lys Gln lle Le	u Leu Asp Ala lle Asp Lys	s Ile Ala		
	200	205	210		
	Asp Asp Trp Asp Asn Arg		Pro Leu Val		
	215	220	225		
	Ala Pro Pro Gln Gly Pro II	e Pro Met Thr Ala Arg Ph			
30	230	235	240		
	Gly Leu Gly Val Pro Arg G	_			
	245	250			

	Asn Cys Val Val Gln Ser Phe Gly Val Ile Gly Gln Ala His Leu			
	260	265	270	
	Glu Leu Pro Arg Pro Asn Lys Arg Ile Arg Asn Gln. Ser Phe Asr			
	275	280	285	
5	Gln Tyr Asn Cys Ser Ile Asn. Asn Lys Thr Glu Leu Glu Thr Trp			
	290	295	300	
	Lys Leu.Val Lys Thr Ser Gly Val Thr Pro Leu Pro. lle Ser Ser			
	305	310	315	
	Glu Ala Asn Thr Gly Leu			
0	320			